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Device for receiving a fluid sample and applications thereof

5 The invention relates to a device for receiving a fluid sample and to the use thereof. The invention relates in particular to a device that makes it possible in particular to sample a small amount of a fluid in a sampling zone and to transport the fluid sampled so as to deposit it, in a depositing zone, on a substrate.

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The device of the invention can also be used as an electrochemical microcell.

15 The invention can be used especially in the biotechnology sector, and in particular in the field, currently in full expansion, of the analysis of biological samples, or in the study of the reactivity or the affinity of a molecule compared with one or more other molecules. The invention can also be used in the
20 more general field of material analysis.

Devices for transferring fluid samples of biological origin, or containing purified molecules produced *in vitro* or *in vivo* are currently of increasing
25 importance. It is known that one of the recent tendencies in this field is to miniaturize the devices and to minimize the amounts of reagents to be used and/or of products to be analyzed or to be studied. In fact, the amounts of products available are often very
30 small, or the products are very expensive. For these reasons, use is increasingly made of devices that make it possible to perform, on appropriate substrates, spot deposits organized in microarrays. These substrates are subsequently brought into contact with known or unknown
35 products that may have an interaction (reactivity or affinity) with respect to the molecule(s) deposited in the microarray. An analysis is then carried out using a

detection system, which may, for example, be optical, chemical, electrochemical, etc.

5 Devices for depositing with or without mechanical contact with the substrate currently exist.

10 In the case of a deposit without contact, the principle is to sample a fluid, containing for example molecules of biological interest, from a sampling zone, and then to place the device above a depositing zone of a substrate, and to deliver a drop of the liquid without the device being in contact with the substrate. Such a device is described, for example, in US patent 5 763 278. The device of this US patent comprises a
15 piezoelectric element which compresses a chamber of small volume and thus ejects a drop of liquid onto substrates of microscope cover select type, or onto any other appropriate support, subsequently allowing analysis using an instrument for demonstrating
20 interactions. Such devices have the advantage of not altering the surface on which the liquid must be deposited, but they are delicate to handle since, in general, the lower end of the device (where the drop to be deposited forms) is very fragile. Such devices also
25 have the drawback of requiring quite large sample volumes. In addition, the complexity of their fabrication and calcination is quite restrictive, especially due to the nature of the materials used, in particular piezoelectric ceramics, which have a
30 tendency to distort over time and with the voltages applied.

35 Among the depositing devices, certain use active fluidic means (pistons, valves, pumps) which have drawbacks both with regard to the complexity of fabrication and the risks of leakages, of blocking of tubing or of formation of bubbles.

As indicated above, devices also exist which operate by contact with the substrate. Some of these devices operate only by virtue of the phenomenon of capillary action. This is the case of the devices described in US
5 patents 5 770 151, 5 807 522 and 6 101 946, which are analyzed hereinafter.

US patent 5 770 151 describes the device for sampling a liquid sample and depositing microdrops of this sample,
10 comprising a hollow tube, one end of which is closed and the other end of which is open. The wall of the tube has, in the region of the open end, a longitudinal gap which promotes the sampling by capillary action of a small amount of liquid when the open end part is
15 immersed in said liquid. Microdrops are subsequently deposited by capillary action, by placing the open end in contact successively with a plurality of spots on a solid surface.

US patent 5 807 522 describes a device for sampling and depositing a liquid sample, comprising two spaced-apart, coextensive members so as to form an elongate capillary channel comprising lateral gaps and ending with a tip. By immersing the tip region in a liquid, a
20 sample is retained in the capillary channel, and when the tip comes into contact with a solid support with sufficient impulsion, the meniscus at the base of the liquid sample is broken, which allows a microdrop of liquid sample to be deposited on the support.
25

US patent 6 101 946 describes a pin for printing a microarray on a support, by depositing microdrops of a liquid sample. This pin comprises a point cut in the shape of a square-based pyramid, comprising a
30 longitudinal gap forming two tips which are brought into closer proximity toward the end of the point. The fabrication of such pins requires high precision machining and is therefore very expensive.
35

All the devices described in the three patents analyzed above comprise a longitudinal gap with the aim of promoting the retention by capillary action of a relatively large amount of liquid. The production of
5 these gaps complicates the fabrication of these devices and requires, in particular for the devices of US patents 5 807 522 and 6 101 946, expensive machining; in addition, these capillary systems become blocked if the sample contains particles in suspension, and they
10 are, moreover, quite difficult to decontaminate.

As will be seen hereinafter, the device of the invention can be used for depositing and fixing on a substrate, in particular by the electrochemical
15 process, a ligand (biological substance or any molecule capable of interaction either with a reagent for analyzing or studying the properties of the ligand, or with a molecule of interest to be detected and/or to be quantified).

20 Patent FR 2 789 401 describes a process for depositing, in an array-like manner, a ligand and electrochemically fixing it on a conducting support. This process can be carried out in particular using a device comprising a
25 conically shaped reservoir made of insulating material (polypropylene) containing a fluid reaction medium and an electrode. The fluid medium contains two types of electropolymerizable monomers, firstly pyrrole, and secondly pyrrole covalently bonded to a ligand. The end
30 of the cone is open and has a small diameter. By bringing this end into contact with a conducting support subjected to an anode voltage relative to the electrode, it is possible to deposit, on the zone of contact with the substrate, a pyrrole polymer, some of
35 the units of which are covalently bonded to the ligand. In such a device, the reservoir contains relatively large amounts of the fluid medium, and the end of the cone has no sampling function, and the reservoir must be filled by means of an active fluidic system (pumps,

valves, etc.). In another embodiment, an electrode in the form of a wire which is immersed in a container containing the fluid reaction medium is used. When the electrode emerges from the fluid, it retains a drop of fluid at its end. The electrode is subsequently brought over the conducting support such that the drop comes into contact with the support while at the same time remaining in contact with the electrode. By applying an appropriate electrical voltage, the formation of a deposit of pyrrole polymer is obtained, as previously. Such a process requires a very precise control of the electrode-support distance. In fact, only the drop, and not the electrode, must come into contact with the support, since a short circuit would prevent the polymerization. It results therefrom that such a process, which is, moreover, relatively non-reproducible, is not suitable for industrial applications. Moreover, the volume of fluid transported by the pin is relatively nonreproducible and subject to drying.

Application WO 00/25925 describes a device for depositing drops of fluid on a substrate. This device comprises a cavity that can communicate with the outside via a capillary channel.

A subject of the present invention is in particular a device for sampling and transporting a fluid, including when the fluid sample which is the source of the fluid to be sampled is available only in very small amounts. This device, which can operate without active fluidic means such as pistons, pumps or valves, is very simple to fabricate and therefore has a relatively low cost price. It does not comprise any fragile elements and can thus be used over a long period of time.

A subject of the invention is a device for receiving, in particular for sampling and transporting, a fluid sample, which comprises an end part having at least one

cavity which opens to the exterior via an opening, said cavity being equipped with a base, characterized in that said end part exhibits a first hydrophobic zone which is adjacent to the cavity opening and a second
5 hydrophilic zone which is adjacent to the first zone and which at least partially covers the base of the cavity such that, when said end part is immersed in said fluid and then emerges therefrom, said cavity retains part of said fluid by means of capillary
10 action.

The receiving device according to the invention is preferably designed so as to form an electrode, in particular a counterelectrode or a working electrode,
15 in an electrochemical cell.

According to the present invention, the term "hydrophobic zone" is intended to mean a zone exhibiting an affinity for a fluid under consideration,
20 in particular a liquid, that is weaker than the hydrophilic zone.

In particular embodiments, the device of the invention can also have the following characteristics, taken
25 alone or, where appropriate, in combination:

- the hydrophobic nature is provided by a hydrophobic coating,
- said hydrophobic coating is deposited on said end
30 part at least at the periphery of said opening (it being understood that this coating must not close up the opening),
- the hydrophobic zone extends into the cavity, optionally to the base thereof, without completely
35 covering the base, and/or extends onto an outer wall of the device,
- the hydrophobic zone is made of an electrically insulating material,

- said hydrophobic coating is made of a material chosen, for example, from Teflon®, such as polytetrafluoroethylene (PTFE), polyvinylidene fluoride (PVDF), perfluoroalkoxy (PFA),
5 homopolymers or copolymers of ethylene, of propylene or of isoprene, polyurethanes and epoxy resins, this list not being limiting,
- the hydrophilic zone is made of a metallic or nonmetallic, electrically conducting material,
- 10 - the end part comprises a body, which is made of an electrically conducting material and/or is coated with an electrically conducting material, the cavity being at least partially formed by this body,
- 15 - said electrically conducting material is chosen in particular from steel, titanium, platinum, gold, silver, graphite and carbon fibers, this list not being limiting,
- said cavity has at least one of the following
20 characteristics:
 - it has a volume sufficient to retain a volume of fluid sample in the range of from 0.1 picoliter to 1 μ l, and in particular from 1 to 50 nl,
 - 25 • it has a depth of 5 μ m to 200 μ m,
 - the cavity depth/opening diameter ratio can vary in the range of from 0.01 to 1, for example from 0.1 to 1,
 - the cavity can have a circular or polygonal
30 transverse cross section,
 - the cavity can have a substantially cylindrical or conical shape, or have a cylindrical wall extended by means of a conical base,
- said device may or may not comprise a damping
35 element for reducing the impacts that may affect said device when it comes into contact via its end part with a solid substrate in order to deposit said fluid sample thereon; said damping element is, for example, a spring,

- said device comprises a rod; the rod can be made of a material capable of elastic deformation, and can comprise at least one part in the shape of an S which plays the role of a damping element,
 - 5 - said device comprises a rod that can slide in another part, in particular a cylinder designed to play the role of a damping element,
 - the hydrophilic nature of the hydrophilic zone can be provided by a coating made of a hydrophilic material.
- 10

Preferably, the cavity for receiving the fluid sample opens directly to the exterior without the involvement of a capillary channel.

15

Thus, the cavity can be emptied and cleaned relatively easily.

According to a particular embodiment, the device of the invention comprises a rod equipped, on the outside or 20 the inside, on the side of the end part, with a sleeve that has a protruding part which extends beyond the end of the rod. The cavity consists, in this case, of the protruding part of the inner wall of the sleeve and of 25 the rod end face. The sleeve is, for example, made of a hydrophobic material. In particular, the rod can be made of a conducting material, and the sleeve of an insulating material, and if it is desired to use the device as an electrode, said rod end face can be 30 polished and/or coated with a relatively unreactive metal, for example platinum or gold, so as to obtain a more stable electrode.

The protruding sleeve can also be made of a conducting 35 material. In this case, at least the end of the protruding part is coated with a layer of hydrophobic material, preferably electrically insulating material. The hydrophobic coating can extend onto the outer wall

of the conducting sleeve and, optionally, onto part of the protruding inner wall.

A subject of the invention is also a process for in particular sampling and transporting a fluid sample using a device as defined above. This process, which can operate without the use of active fluidic means, comprises the steps consisting in:

- a) immersing the end part comprising said cavity in a container containing a fluid to be sampled, and then removing it therefrom, and
- b) bringing said end part into contact with a solid substrate.

According to particular embodiments:

- the end part is subsequently moved away from the substrate, so as to leave, as a deposit on the substrate, a drop of the fluid sample,
- if desired, steps a) and b) are repeated as many times as necessary for depositing a plurality of identical or different fluid samples on the solid substrate, so as to form, on said substrate, deposits in the form of a matrix array. When the samples are different, a rinsing-drying operation will be necessary.

This process can be used in particular with a fluid sample which contains biological molecules or substances to be deposited and/or to be immobilized on the substrate. It can also be used for transporting a fluid to another fluid solution, so as to produce a dilution, for example.

The process of the invention also makes it possible to use the receiving device as an electrode. For this, the device comprises a body made of a conducting material, and said end part is equipped with an insulating and hydrophobic coating or sleeve which, of course, does not close up the cavity opening. The substrate is made

of a conducting material or contains one or more
conducting zones and, after said step of bringing into
contact, the assembly forms an electrochemical cell
comprising at least two independent electrodes. One or
5 more additional electrode(s) can be added either to the
device, or to the substrate.

The fluid can comprise an electrolyte and, optionally,
other compounds in suspension.

10

The process can comprise the step consisting in
carrying out an electrochemical-type analysis of the
solution or suspension sampled.

15 The process can comprise the step consisting in using
the abovementioned assembly as an electrochemical cell
and passing an electric current, or simply measuring a
potential difference, between said end part and said
substrate or between the end part and a conducting zone
20 of the substrate, by means of the sample, containing an
electrolyte, which is in contact both with the
conducting body and with the substrate. Using a
galvanostatic or potentiostatic assembly, it is
possible to determine the current and potential
25 characteristics of the sampled fluid sample to be
analyzed or of the substrate, without any notable
modification of the composition of the sample, since
the concentrations of the electroactive substances
dissolved are virtually unmodified by the measurements
30 carried out.

Using, for example, the device of the invention as a
working electrode and the conducting support, in
particular a metal strip, as a counterelectrode, the
35 cavity of the device constitutes an electrolyte
microcell for in particular studying the reactions
which occur at the working electrode. Such a device
makes it possible to always have the same distance
between the working electrode and the counterelectrode.

It is also possible to use the process of the invention for immobilizing one or more biological molecules or substances on the conducting substrate according to an electrochemical method of electrodeposition. In this case, the substrate constitutes the working electrode and the end of the receiving device serves as a counterelectrode.

Such a process of electrodeposition can be carried out in particular when the fluid contains a monomer that is electropolymerizable, for example by anodic oxidation. The electric current is then passed between the body and the substrate, bringing said substrate to a potential required for polymer formation. Thus, the end of the sampling device, made of a conducting material, plays the role of a counterelectrode, such that the monomer will polymerize in contact with the conducting substrate, by anodic oxidation, and form a point deposit, also called spot, that adheres on said substrate. Such a process therefore makes it possible to produce microspots of polymer, optionally arranged as an array, on a conducting surface.

A subject of the invention is also a process for forming an electrochemical cell, the process comprising the following steps:

- providing a receiving device which comprises an end part having at least one cavity which opens to the exterior via an opening, said cavity being equipped with a base, this end part exhibiting a first electrically insulating hydrophobic zone which is adjacent to the cavity opening and a second electrically conducting hydrophilic zone which is adjacent to the first zone and which at least partially covers the base of the cavity,
- providing a receiving surface, in particular a substrate, having at least one conducting zone,

- sampling a fluid sample by means of the receiving device,
 - bringing the end part of the receiving device into contact with the conducting zone of the receiving surface, the first hydrophobic zone being designed so as to electrically insulate the second conducting hydrophilic zone from the conducting zone of the receiving surface.
- 10 A subject of the invention is also a process comprising the following steps:
- providing a receiving device which comprises an end part having at least one cavity which opens to the exterior via an opening, said cavity being equipped with a base, this end part exhibiting a first electrically insulating hydrophobic zone which is adjacent to the cavity opening and a second electrically conducting hydrophilic zone which is adjacent to the first zone and which at least partially covers the base of the cavity,
 - providing a receiving surface, in particular a substrate, having at least one conducting zone,
 - sampling a fluid sample by means of the receiving device,
 - bringing the end part of the receiving device into contact with the conducting zone of the receiving surface, the first hydrophobic zone being designed so as to electrically insulate the second conducting hydrophilic zone from the conducting zone of the receiving surface,
 - establishing an electric current between the hydrophilic zone of the receiving device and the conducting zone of the substrate or measuring an electrical parameter, for example a potential difference, between the conducting zone of the receiving device and the conducting zone of the receiving support.

The process can comprise the following step:

- establishing an electric current, in particular a pulsed current, between the hydrophilic zone of the receiving device and the conducting zone of the substrate in order to polymerize a substance,
5 contained in the cavity of the receiving device.

The process can, as a variant, comprise the following steps:

- measuring an electrical parameter, in particular a
10 potential difference, between the conducting zone of the receiving device and the conducting surface, for example a steel sheet,
- repeating the preceding step in order to carry
15 out, for the conducting surface, a mapping relating to a physical or chemical characteristic, for example an oxidation state, using the measurements obtained.

Particular embodiments of the invention will now be
20 described in greater detail, by way of illustration, with reference being made to the attached drawings in which:

- figures 1 to 7 represent, diagrammatically and
25 partially, particular embodiments of the end part of the device of the invention,
- figures 8 and 9 represent, diagrammatically and partially, embodiments of a receiving device with a damper,
- figure 10 represents, diagrammatically and
30 partially, a support of a counterelectrode in accordance with the invention,
- figures 11 and 12 represent, diagrammatically and partially, an indicating electrode in accordance with two examples of implementation of the
35 invention, and
- figures 13 to 15 illustrate another example of implementation of the invention.

Represented in figure 1, very diagrammatically, is a receiving device 1 in accordance with the invention.

5 The device 1 comprises a rod 2, an end part 2' of which is provided with a receiving cavity 3.

In the example considered, the cavity 3 is in the shape of a cylinder having an axis X parallel to the rod 2, with an inner wall 4 and a base 5.

10

The rod 2 has, at its end, a section 6 covered with a hydrophobic coating 8.

15 In the example considered, the rod 2 is made of a conducting material exhibiting hydrophilic properties, it being possible for this conducting material to be, for example, gold, platinum or a stainless steel of the 316L stainless steel type.

20 The section 6 extends to the periphery of the opening 7 of the cavity 3.

In the example of figure 1, the coating 8 extends only over the section 6, without going over into the cavity 25 3, nor onto the outer wall 10 of the rod 2.

As a variant, the coating 8 can extend, as illustrated in figure 2, into the cavity 3, partially covering the inner wall 4.

30

This coating 8 may or may not reach the base 5.

The coating 8 can also extend onto the outer wall 10 of the rod 2.

35

In the examples which have just been described, the receiving cavity 3 is produced in a hollowing-out of the rod itself.

Represented in figure 3 is a receiving device 15 in accordance with another example of implementation of the invention, in which the receiving cavity 16 is formed by a sleeve 17 fitted at one end of a rod 18.

5

The cavity 16 has a base defined by the section 19 of the rod 18.

The sleeve 17 comprises a first part 17a engaged on the rod 18 and a second part 17b protruding from the section 19.

The sleeve 17 is made of a hydrophobic material, consisting, for example, of a heat-shrinkable sheath made of plastic.

15

In the example of implementation illustrated in figure 4, the rod 18' comprises an annular necking 20 on which the sleeve 17 is fitted.

20

Represented in figure 5 is a receiving device that differs from that described with reference to figure 1 by virtue of the fact that the end of the rod 2' is at least partially beveled, it being, for example, semi-beveled or completely beveled.

25

In the example of implementation illustrated in figure 6, the receiving device 20 comprises a rod 21 at one end of which is attached a metal insert 22 comprising a receiving cavity 23.

30

The insert 22 comprises a cylindrical outer wall 24 covered with a hydrophobic coating 25.

Represented in figure 7 is a receiving device 35 with a metal rod 36 having at one end a head 37, part of the rod 36 and this head 37 being embedded in a coating made of a hydrophobic material 38.

35

This coating 38 comprises, at right angles with the head 37, a cavity 39 for receiving a fluid sample.

5 Represented in figure 8 is a device 30 in accordance with the invention, comprising a metal rod 31, which has a part 32 folded in the shape of an S, designed so as to define an elastically deformable zone forming a damper.

10 This damper can thus be produced in a particularly simple manner and makes it possible to damp impacts along a direction perpendicular to the plane of the substrate 33. In the example considered, the substrate 33 comprises a gold plate.

15 The lower end 34 of the rod 31 defines a receiving device 20 described with reference to figure 6.

20 In the example of implementation illustrated in figure 9, the receiving device is attached to a rod 41 which has, at an upper end, a head 42 which fits into a housing of a support 40.

25 This head 42 is returned to its starting position by means of a spring 43 distinct from the rod 41.

30 Thus, the rod 41 can be free of a part folded in the shape of an S such as that described with reference to figure 8.

The spring 43 can be replaced by any other element with an elastic return, such as an elastomeric material, for example.

35 This support 40 can be attached to a manipulating arm of an automated device for moving the rod along horizontal and vertical directions.

Such an automated device can be designed so as to be able to actuate a plurality of receiving devices.

Using an automated 3-axis device, it is possible to
5 deposit drops on a substrate according to a matrix array.

A typical method of use consists in bringing the receiving device over the sampling zone, and moving the
10 rod vertically downward until its end is immersed in the fluid to be transferred. This is followed by a horizontal movement until it is plumb with the depositing zone on a substrate, and a vertical descent until there is contact with the substrate, and
15 deposition of a microdrop by capillary action. Next, the device is raised back up vertically, and then moved to a zone for cleaning the end, for example by water jet, and then air jet for drying. The operations can then be repeated with the same fluid sample or another
20 fluid sample.

When the hydrophobic coating or sleeve is made of an insulating material, it is possible to use the device of the invention as an electrode, the cavity 23 playing
25 the role of an electrochemical microcell.

Various applications of the invention will now be described in greater detail hereinafter.

30 **Example 1: Production of protein chips by electrochemical deposition**

This involves producing a chip comprising 60 spots of 6 different molecules, 10 copies of each being
35 immobilized; each spot is arranged on a virtual square grid pattern of 8×8 spots, with a gap of $700 \mu\text{m}$ between the center of each spot and on a total surface area of $5 \times 5 \text{ mm}^2$. Four zones will not be

functionalized with biological species, the substrate will remain "bare".

5 The six different molecules are antibodies: an anti-hCG antibody (Sigma), an anti-peptide mAb 11E12 (Sanofi Diagnostics Pasteur), an anti-HSA antibody (Sigma), an anti-avidin antibody (Sigma), an anti-rabbit IgG antibody (Sigma) and an anti-BSA antibody (Sigma).

10 The final objective of the experiment is to observe the interactions in parallel, in real time and without a label, of these antibodies with the molecules against which they are directed, injected successively in contact with the chip, by means of the Surface Plasmon
15 Resonance imaging technique such as that described in application WO 02/48689. All these molecules are coupled beforehand to pyrrole monomers on their NH₂d bond. After this, each protein coupled to one or more pyrrole molecules, at a concentration of 10 µM in a
20 reaction medium consisting of 50 mM NaH₂PO₄ (Sigma) + 50 mM NaCl (Merck), at a pH of 6.8, is placed at the bottom of one of the wells of a conical-bottom 96-well microplate. A few microliters of product, typically less than or equal to 5 µl, are sufficient to make it
25 possible to carry out several tens of deposits per species.

The substrates used in this example are prismatic substrates, with a base of 12.5 × 25 mm² and a height
30 of 9 mm (made of BK7 or SF11 glass), onto which have been deposited a layer of chromium of approximately 20 angstroms, which serves as an attachment layer, and a layer of gold of approximately 500 angstroms (depositions carried out by evaporation under vacuum).
35 Substrates of these types are particularly suitable for measurements made by Surface Plasmon Resonance.

The layer of gold is then connected to an EGG 273-type potentiostat forming a working electrode outlet. As

regards the counterelectrode part of this electrochemical cell, the procedure is as follows:

- a receiving device, for example the receiving device 30 described with reference to figure 8, is inserted into a stainless steel cylinder 50 with a hollowing-out 51 receiving the device 30, as illustrated in figure 10,
- the cavity 23 has a circular cross section, with an internal diameter of 250 μm ; the external diameter of the insulating sleeve 25 is 450 μm and the depth of the cavity 23 is 50 μm , which corresponds to a total cavity volume of approximately 2.5 nl,
- to maintain the device 30 in the hollowing-out 51, a stainless steel blocking screw 52 is used, which also makes it possible to establish the electrical connection between the conducting part of the cavity 23 of the device 30 and the counterelectrode outlet, via an electrical wire 53,
- the cylinder 50 can be maintained in the vertical position either in a chuck, or can be installed on a manipulating arm 54, represented very diagrammatically as dashed lines in figure 10, for example of an industrial automated device with 3-axis movement, called GENESIS sold by the company TECAN for example, by virtue of threading cuts on the upper part of this cylinder; it is noted that a reference electrode is of no use in this present case, the latter being directly connected to the counterelectrode.

The manipulating arm 54 holding the cylinder 50 is placed vertically to the well containing the anti-hCG antibody.

The manipulating arm 54 descends into the well such that the cavity 23 of the device 30 is completely immersed in the solution.

A mechanical contact at the bottom of the well is possible and does not impair the functionality of the device. Some of the solution, a few nl in this case,
5 penetrates into the cavity by capillary action.

The manipulating arm 54 is moved back up vertically and is displaced above the depositing zone on the gilded prismatic substrate, and more particularly above one of
10 the predetermined zones of the array. The manipulating arm 54 then descends until mechanical contact is obtained between the device 30 and the substrate.

The electrical contact, which is made between the
15 conducting base of the cavity 23 and the substrate via the conducting reaction medium, does not necessarily require the receiving device 30 to be in mechanical contact with the substrate. It is, however, preferable to effect such a mechanical contact.

20 Once the arm 54 is immobilized, a potential difference of + 2.4 V is established for 250 ms between the counterelectrode and the working electrode by virtue of the EGG 273 potentiostat. There is then formation of a
25 thin film of polypyrrole on the substrate, by which the biomolecules, i.e. the anti-hCG antibodies, are attached to the gold-coated prismatic support.

The manipulating arm 54 can then be moved back up and
30 put back into the preceding well in order to take another sample; the rinsing and the drying of the device are not essential in this case since the same product is sampled several times.

35 Once the ten spots have been formed according to the same process, the manipulating arm 54 is moved to be vertical to a well of the microplate filled with ultrapure water. The arm 54 is then moved in and out of this well three times so as to correctly rinse the

device 30, which, without distinction, may or may not come into contact with the bottom of the well without impairing the future functionality thereof. The manipulating arm 54 is then brought into contact with an absorbent paper, for example an optical paper sold by the company Kodak. This drying operation is carried out three times, at three different places on the absorbent paper.

After this drying phase, the manipulating arm 54 is controlled with a view to sampling a second antibody, an anti-HSA, for example, according to the sequence described above, so as to deposit the 10 spots electrochemically. The process is carried out in this way for the other four species.

Similarly, 96 spots of different species can be deposited, by performing a cleaning-rinsing-drying phase at the end of each electrodeposition.

20

Example 2: Production of a 384-spot chip exhibiting DNA sequences relevant for studying cystic fibrosis
(deposition by passive adsorption)

In this example, the substrates are microscope slides ($75 \times 25 \times 1 \text{ mm}^3$, sold under the name ESCO by the company VWR International) on which are deposited beforehand a layer of chromium of approximately 20 angstroms and a layer of gold of approximately 500 angstroms (depositions carried out by vacuum evaporation).

This slide is functionalized with a coating which promotes the immobilization of biomolecules by electrostatic interactions. It is a monolayer of 11-mercaptoundecanoic acid (MUA) deposited onto the gold and then a monolayer of polyethyleneimine (PEI) (method described by Bassil et al., Sensors and Actuators B94 (2003) 313-324). This surface is then

brought into contact with a solution of extravidin (Sigma) at 0.2 g/l in PBS (Sigma) for 30 minutes before being rinsed with water. The extravidin then attaches to the PEI by means of the electrostatic interactions.

5

The substrate is placed in the working zone of a 3-axis automated device, for example that sold under the name Q-Array by the company Genetix, which already has predetermined positions for microscope slides of this format and also for standard microplates with a support comprising an integrated damping device, the damping occurring under its own weight, and into which the device 40 described, for example, with reference to figure 9 is inserted.

15

The dimensions of the device are, in this case, as follows: the internal diameter of the sample-receiving cavity is 100 μm , the depth thereof is 50 μm and the external insulating PTFE sleeve diameter is 300 μm .

20

Several oligonucleotide sequences (300 different sequences in total), functionalized with a biotin in the 5' position, are placed separately in the wells of a 384-well microplate, in a PBS buffer, in the presence of 1.5 M of betaine so as to prevent the species from drying out too rapidly on the chip. The concentration of the sequences is 1 μM in each of the wells. These sequences were chosen so as to determine with certainty the type of mutation implicated in cystic fibrosis.

25

30 Three copies of each species are deposited, distributed randomly over a virtual rectangular array composed of 16 \times 64 points 400 μm apart (1024 measuring points in all).

35

The arm which carries the device previously described moves the rod down into one of the wells of the microplate. The product is sampled by capillary action when the rod is immersed in the liquid containing the oligonucleotides.

The manipulating arm 54 then moves back up and is positioned above one of the points of the array. The arm descends vertically and, when there is mechanical
5 contact between the substrate and the device, the latter deposits on the substrate a part of the volume sampled in the form of a microdrop having a volume of approximately 1 to 2 nl.

10 The arm 54 then returns above the same well as previously, and carries out the previous cycle a further two times in order to produce two other spots of the same biological species.

15 Once the three spots of each of the species have been spotted, the arm again moves back up and is placed vertically to a fountain spraying ultrapure water on the rod, so as to remove the fluid still present in the cavity or on the outer wall of the device.

20 The arm subsequently places the device over a drying element producing, for example, a stream of hot dry air and remains there for a few tens of seconds.

25 The device is then ready to sample a further product from another well, until all the types of oligonucleotide sequences have been sampled and deposited.

30 **Example 3: Fluorescence techniques**

Finally, the analysis of the chip can be carried out by fluorescence techniques. The aim is to compare the expression profile of an affected patient with respect
35 to a normal patient. To do this, the DNA of a normal patient is labeled beforehand with a fluorescent label (Cy3, for example, Sigma) and that of an affected patient is labeled with another fluorescent label (Cy5, for example, Sigma). The sera of the two patients are

mixed and this mixture is brought into contact with the functionalized chip. The products are left in contact for 30 minutes at 37°C. The chip is then rinsed and inserted into a fluorescence reader, for example the
5 Genechip™ Scanner 3000. The analysis of the fluorescence of the two labels on each of the spots, corresponding to the various oligonucleotide sequences, then makes it possible to determine which are the genotypes overexpressed or underexpressed in the
10 affected patient compared with the normal patient.

Example 4: Parallel deposition

The invention can be implemented for parallel
15 deposition, with or without electrochemistry, with 8 rods which sample, from a 1536-well plate, around 8 different substrates (for example, 8 rods installed on an automated device called GENESIS sold by the company TECAN).

20

Example 5: Use of the receiving device as an indicating electrode or working electrode

In the following two examples, the rod is used as a
25 working electrode in a two-electrode electrochemical microcell. This type of device makes it possible, for example, to characterize molecules in the reduced or oxidized state or to study the synthesis of polymers by electrochemistry.

30

Example 6: Use in galvanostatic mode

Using an electrode with a very small surface area, of the order of 1 mm², which is called an indicating
35 electrode, it is possible to determine current-potential characteristics while at the same time conserving the system virtually without modification of composition, i.e. without substantially modifying the concentrations of the electroactive substances

dissolved in the electrolyte, despite the passage of the current.

5 A solid electrode 60 illustrated in figure 11 is produced by inserting a rod 61 made of platinum, gold, silver, graphite or stainless steel, with a diameter of between 0.5 mm and 2 mm, into an insulating sheath 62 made of insulating glass, polyethylene or Teflon® for example, and releasing the straight section of the rod
10 so as to bring it into contact with the solution. A planar disk electrode is thus obtained. The end of the rod in contact with the solution can be polished with, for example, diamond paste.

15 The cavity 63, having a continuous wall, makes it possible to create an electrochemical microcell filled, by capillary action, with the sample to be analyzed.

The working electrode 60 formed by the rod 61 is
20 connected to the outlet of a working electrode of a potentiostat. This connection can be made directly on the rod or on a metal piece into which the rod fits, and designed so as to adapt to an automated device.

25 The counterelectrode 65 may be a sheet of platinum, a gold strip, a plastic support coated with ITO (indium tin oxide), or a silicon plate, for example.

The reaction medium may be an ionic solution based on
30 Li^+ , ClO_4^- ions or PBS for example, containing the chemical species to be analyzed.

The end of the electrode 60 is brought into contact with the counterelectrode 61 and a current of a few
35 tens of microamps is applied. The voltage is then measured.

This device can, similarly, be used in the potentiostatic mode.

In this case, a voltage is applied between the two electrodes and the current generated by this voltage is analyzed.

5

As previously, the rod is used as a working electrode, a gold strip is used as a counterelectrode and the cavity is used as an electrochemical microcell. The reactions which take place at the working electrode consisting of the rod are then studied.

10

This device makes it possible to always have the same distance between the working electrode and the counterelectrode.

15

The sleeve 62 can either be of insulating nature, or of conducting nature and coated with a layer 66 of an insulating material 67, for example a rigid insulating Teflon®, as illustrated in figure 12.

20

Example 7: Use of the rod as an auxiliary electrode (counterelectrode)

In this configuration, the function of the rod will be to serve as a counterelectrode and a microcell. This makes it possible to produce microspots of polymer on a metal surface of a few hundred μm in diameter.

25

The rod is made of stainless steel or of stainless steel coated with a metal, for instance platinum, gold or silver. The sleeve is made of stainless steel possibly coated with a metal as regards its internal part and coated with Teflon® on its external part. The sleeve can also consist of an insulating material.

30

35

A voltage of approximately 2 V is applied using a potentiostat or a voltage generator for example, between the rod and the gold strip which serves, in the example under consideration, as a working electrode.

The cavity is filled with an ionic solution containing, for example, pyrrole and the rod then comes into contact with a glass slide coated with chromium and gold thus forming an electrochemical microcell. A potential is then applied between the two electrodes. The current and the charge for synthesis of the polymer (polypyrrole) thus formed, on the surface of the gold-coated slide, are recorded. Several spots can thus be produced on the same surface.

The damping system makes it possible not to damage the rod and the gold strip. The gold strip is also protected by the layer of "soft" Teflon® at the end of the rod.

Coating the inside of the cavity with a metal such as platinum can make it possible to improve the electrochemical synthesis of the polymer.

Example 8: Synthesis of deposits

Use is, for example, made of a receiving device 70 comprising a rod 71 made of 304L stainless steel (surgical quality) and a sleeve 72 made of Teflon®, as illustrated in figures 13 and 14.

The Teflon® sleeve 72 protrudes below the end of the rod 71 so as to define a cavity 73.

This cavity 73 has, for example, a diameter of approximately 260 μm and a depth of approximately 100 μm and makes it possible to receive a solution to be polymerized 74.

The stainless steel rod 71 serves as a counterelectrode.

The solution to be polymerized 74 is deposited on a substrate 75 coated with a layer of gold serving as a working electrode, as illustrated in figure 15.

5 The rod 71 and the substrate 75 are connected to a potentiostat 76.

The synthesis of deposits is carried out by the electrosputting method by applying an electric pulse
10 through the rod 71.

The Teflon® sleeve 72 makes it possible to insulate the counterelectrode from the working electrode, the cavity 73 forming an electrochemical cell in which the
15 electric pulse triggers the polymerization of the solution.

The receiving device 70 also makes it possible to capture a drop of approximately 50 nl by immersing said
20 receiving device in the solution to be polymerized, and to ensure its transport to above the working electrode.

The rod 71 is placed on a supporting structure (not represented) in which it can slide vertically under the
25 action of its own weight. The movements are provided by motorized jackscrews controlled by an automated device.

The conditions for the electrosputting (potential, time) are optimized so as to obtain deposits of pyrrole
30 and of pyrrole ODN. During the polymerization, the charge delivered by the potentiostat 76 is recorded in the form of a chronoamperogram.

Once the deposits have been made, a hybridization is
35 carried out with a labeled complementary ODN in order to demonstrate the spots containing ODNs. The detection is in this case carried out using a fluorescence microscope equipped with a black and white CCD camera

for image acquisition. The fluorescence intensities are expressed as levels of gray.

**Example 9: Process for carrying out redox mapping of a
5 conducting surface**

This example concerns the use of a receiving device according to the invention for characterizing a metal surface, such as a steel sheet, and performing two-
10 dimensional mapping of the oxidation state thereof.

The receiving device used in this case is substantially the same as that used in the preceding example.

15 A layer of silver is added by electrochemical reaction at its hydrophilic end, i.e. on the cavity base.

The electrolyte used for measuring the residual potential between the silver-coated electrode and the
20 metal sheet is, for example, 100 mM KCl. The receiving device and the electrolyte are deposited on a first point of the surface to be mapped, and the value of the potential difference between the metal sheet and the receiving device is recorded. The receiving device is
25 then rinsed, dried and filled again with electrolyte, followed by deposition at a second point of the surface to be mapped.

This process makes it possible to detect any points of
30 oxidation of the steel. The various treatments of the metal sheet can therefore be readily studied and compared.